NOV 21 2008

510(k) Summary
510(k) Number K082604

Viztek, Inc.
6491 Powers Avenue
Jacksonville, FL 32217
Phone: 800.366.5343
Fax: 904.448.9936
Date Prepared: July 30, 2008
Contact: Bruce Ashby, Sales and Marketing Manager

1. **Identification of the Device:**
   Proprietary-Trade Name: Viztek DR Series Digital Diagnostic Digital X-Ray Systems (Multiple Models)
   Classification Name: Stationary x-ray system, Product Code 90 KPR and Solid State X-Ray Imager (Flat Panel/Digital Imager) 90 MQB,
   Common/Usual Name: Digital Stationary Diagnostic X-Ray System

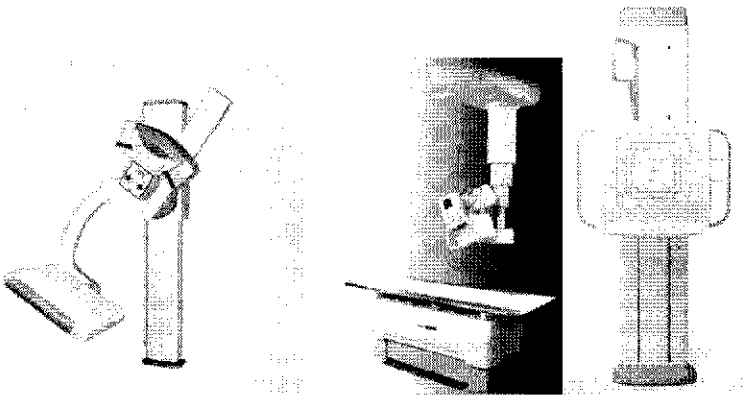
2. **Equivalent legally marketed device**: URS LP Digital, Millennium, Radpro (Sedecal) K042876, and SmartRad (CMT Medical Technologies) K003438.

3. **Indications for Use (intended use)** This digital radiographic system is intended for use by a qualified/trained doctor or technician on both adult and pediatric subjects for taking diagnostic radiographic exposures of the skull, spinal column, chest, abdomen, extremities, and other body parts. Applications can be performed with the patient sitting, standing, or lying in the prone or supine position.

4. **Description of the Device**: This digital diagnostic x-ray system consists of a tubehead/collimator assembly mounted on a ceiling suspension OR a U-Arm, along with a generator, generator control, and an elevating x-ray table. Power ratings for the available generators are in the rage of 50 kw to 80 kW. Exposure voltage range varies from 40 125 KV or 40 150 kV with current of 300 - 100 mA. Exposure time is 1 ms – 10 s. Models: Model DR3000 (U-arm single detector) and Model DR4000 (dual detector-ceiling suspension and table and upright bucky) and DR1000 single detector (standing bucky and ceiling suspension. The digital subsystem is the SmartRad (CMT Medical Technologies) K003438.

5. **Safety and Effectiveness, comparison to predicate device**. The results of bench and test laboratory indicates that the new device is as safe and effective as the predicate devices.

Model 3000 **Models 4000 and 1000**



6. Substantial Equivalence Chart

Characteristic	URS LP Digital, Millennium, Radpro (Sedecal) K042876	Viztek DR Series
Intended Use:	Intended for use by a qualified/trained doctor or technician on both adult and pediatric subjects for taking diagnostic radiographic exposures of the skull, spinal column, chest, abdomen, extremities, and other body parts. Applications can be performed with the patient sitting, standing, or lying in the prone or supine position.	SAME
Configuration	U-Arm mount or Ceiling Suspension	SAME
Performance Standard	21 CFR 1020.30	SAME
Generator	High frequency made by Sedecal	Uses same generator made by Sedecal
Digital Panel	CANON CXDI 50G	Pixium 4600
Number of panels	One	One or two
Electrical safety	Electrical Safety per IEC-60601. UL listed	SAME

7. Conclusion

After analyzing bench and external laboratory testing to applicable standards, it is the conclusion of Viztek Inc that the Viztek DR Digital Diagnostic X-Ray Systems are as safe and effective as the predicate device, have few technological differences, and has no new indications for use, thus rendering them substantially equivalent to the predicate devices.

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

NOV 21 2008

VIZTEK
% Mr. Daniel Kamm, P.E.
Principal Consultant
Kamm & Associates
PO Box 7007
DEERFIELD IL 60015

Re: K082604
Trade/Device Name: Viztek DR Series Diagnostic Digital X-Ray Systems (Multiple Models)
Regulation Number: 21 CFR 892.1680
Regulation Name: Stationary x-ray system
Regulatory Class: II
Product Code: KPR
Dated: September 1, 2008
Received: September 11, 2008

Dear Mr. Kamm:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

This letter will allow you to begin marketing your device as described in your Section 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Office of Compliance at one of the following numbers, based on the regulation number at the top of this letter:

21 CFR 876.xxx	(Gastroenterology/Renal/Urology	240-276-0115
21 CFR 884.xxx	(Obstetrics/Gynecology)	240-276-0115
21 CFR 894.xxx	(Radiology)	240-276-0120
Other		240-276-0100

Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). For questions regarding postmarket surveillance, please contact CDRH's Office of Surveillance and Biometrics' (OSB's) Division of Postmarket Surveillance at 240-276-3474. For questions regarding the reporting of device adverse events (Medical Device Reporting (MDR)), please contact the Division of Surveillance Systems at 240-276-3464. You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (240) 276-3150 or at its Internet address http://www.fda.gov/cdrh/industry/support/index.html.

Sincerely yours,

Joyce M. Whang, Ph.D.
Acting Director, Division of Reproductive,
Abdominal, and Radiological Devices
Office of Device Evaluation
Center for Devices and Radiological Health

Enclosure

Indications for Use

510(k) Number (if known): K082604

Device Name: Viztek DR Series Diagnostic Digital X-Ray Systems (Multiple Models)

Indications For Use:
This Digital Radiographic System is intended for use by a qualified/trained doctor or technician on both adult and pediatric subjects for taking diagnostic radiographic exposures of the skull, spinal column, chest, abdomen, extremities, and other body parts. Applications can be performed with the patient sitting, standing, or lying in the prone or supine position.

Prescription Use X AND/OR Over-The-Counter Use ____.
(Part 21 CFR 801 Subpart D) (21 CFR 807 Subpart C)

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

(Division Sign-Off)
Division of Reproductive, Abdominal, and Radiological Devices
510(k) Number K082604